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FDA has “gone rogue” by approving new COVID booster shots without clinical trials, warns Dr. Robert Malone

09/15/2023 // Ethan Huff // [2.2K](#) Views

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Without any clinical trials whatsoever to back their alleged safety and efficacy, the all-new Wuhan coronavirus (COVID-19) "booster" shots now being pushed by the Biden regime received the green

light from the *U.S. Food and Drug Administration* (FDA) this week, prompting Dr. Robert Malone to [state that](#) the corrupt federal agency has "gone rogue."

In an appearance on EpochTV's "Crossroads" program on September 11, the day the FDA approved the new booster shots, Dr. Malone tore apart the corrupt federal agency for abandoning all remaining semblances of honesty and integrity.

"It's difficult to conclude anything other than the FDA is no longer feeling bound by their own rules and regulations," Dr. Malone said. "The term is: they've gone rogue."

Attesting to the lack of clinical trial data, particularly in humans, the all-new COVID booster shots never should have received approval and emergency use authorization (EUA) from the FDA in the first place.

The FDA's argument in support for the shots claims that it already has all the evidence it needs without clinical trials because "neutralizing antibodies as detected in mice and their cross-reactivity are correlative protection."

According to Dr. Malone, that claim "is a lie" as "there are no established correlates of protection for SARS-CoV-2."

(Related: Earlier last year, Dr. Malone [confirmed that](#) life insurance company death claims are up bigtime ever since the launch of Operation Warp Speed.)

FDA says it relied on "manufacturing data" from Pfizer, Moderna to approve their mRNA COVID booster "vaccines"

In a statement to go along with its approval and EUA notice, the FDA said that a full evaluation of the "manufacturing data," meaning the data that Pfizer and Moderna provided to the FDA, shows that the shots are safe and effective – and this is enough for the FDA.



The FDA also cited "non-clinical immune response data on the updated formulations including the XBB.1.5 component" as evidence that the shots are safe and effective for human use.

Because the risk-benefit profile for all the other FDA-approved and FDA-authorized COVID "vaccines" is already "well understood," according to the agency, there is no reason to continue testing all the latest booster shots because they are "a good match for protecting against the currently circulating COVID-19 variants."

On September 12, the *U.S. Centers for Disease Control and Prevention* (CDC) followed through with issuing its own recommendation in favor of the shots, urging all Americans to rush out and get jabbed as soon as possible.

This recommendation does not sit well with Florida Surgeon General Dr. Joseph Ladapo, who joined Dr. Malone in stating that "there's essentially no data" to back the new shots.

"Not only that, but there are a lot of red flags," he added, noting that studies show the shots produce *negative* efficacy over time.

"There's been no clinical trial done in human beings showing that it benefits people, there's been no clinical trial showing that it is a safe product for people," he further said.

What is work, earlier iterations of these shots are linked to serious cardiac problems such as heart inflammation – and the new ones more than likely do the same.

"It's truly irresponsible for FDA, CDC, and others to be championing something ... when we don't know the implications of it," he further stated.

Dr. Malone also commented on the negative efficacy aspect of the shots, explaining that not only is there no benefit to getting jabbed, but that it is also *harmful* to get jabbed. In other words, a person is better off relying on natural immunity without any vaccines, period.

Once the new COVID shots start getting injected into people's arms in the coming weeks, expect more serious injury and death reports to emerge. Learn more at [Vaccines.news](https://vaccines.news).

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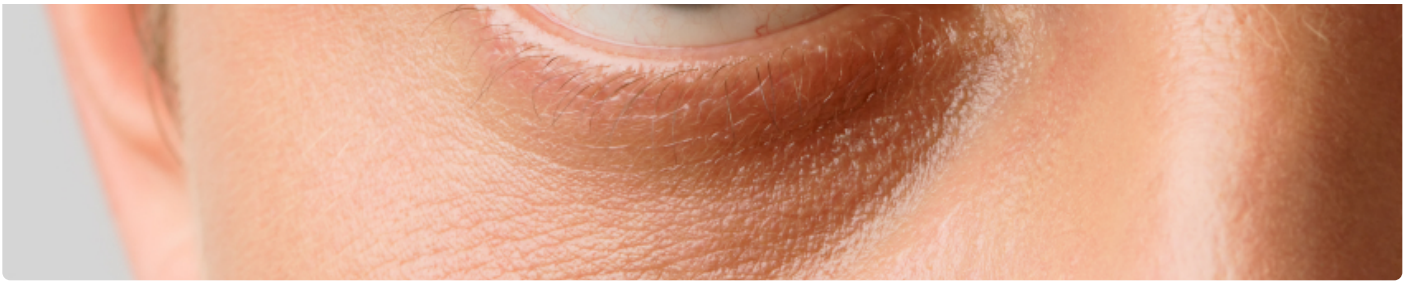
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